Docket No.: 20605/1203866-US1 (PATENT)

## IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

	A. Rizzieri et al.	
Appli	cation No.: 10/008,062	Confirmation No.: 5735
Filed:	October 19, 2001	Art Unit: 1642
For:	ANTI-TENASCIN MONOCLONAL ANTIBODY THERAPY FOR LYMPHOMA	Examiner: Alana M. Harris, Ph.D.

## INFORMATION DISCLOSURE STATEMENT (IDS)

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

In ... Detect Acceliantian of

Dear Sir:

This Information Disclosure Statement is submitted in accordance with 37 C.F.R. 197, 198, and it is requested that the information set forth in this statement and in the listed documents be considered during the pendency of the above-identified application, and any other application relying on the filing date of the above-identified application or cross-referencing it as a related application.

	<ol> <li>This IDS should be considered, in accordance with 37 C.F.R. 1.97, as it is filed: of the boxes A-D)</li> </ol>
A.	within three months of the filing date of the above-identified national application of within three months of the entry into the national stage of the above identified national application
В.	before the mailing date of a first office action on the merits, or a first office action after filing a request for continued examination.
x C.	after (A) and (B) above, but before final rejection or allowance, and Applicants have made the necessary statement in box "i" below or paid the necessary fee in box "i" below.

(check one of the boxes "i" and "ii" below:)
i. Counsel states that, upon information and belief, each item of information listed herein was (check one of boxes (a) or (b))
<ul> <li>(a) first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of this IDS; or</li> </ul>
(b) not cited in a communication from a foreign patent office in a counterpart foreign application and, to the knowledge of undersigned after making reasonable inquiry, was not known to any individual designated in 1.56(c) more than three months prior to the filing of this IDS.
x ii. Payment for the fee set forth in 1. 17(p), presently believed to be \$180, is enclosed.
D. after (A), (B) and (C) above, but before payment of the issue fee: Applicant petitions under 37 C.F.R. 1.97(d) for the consideration of this IDS. Under 37 C.F.R. 1.17(f) a check in the amount of \$180.00 is enclosed. Counsel certifies that, upon information and belief, each item of information listed herein was
(check one of the boxes "a" and "b" below:)
<ul> <li>(a) first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of this IDS; or</li> </ul>
(b) was not cited in a communication from a foreign patent office in a counterpart foreign application and, to the knowledge of undersigned after making reasonable inquiry, was not known to any individual designated in 1.56(c) more than three months prior to the filing of this IDS.

2. In accordance with 37 C.F.R. 1.98, this IDS includes a list (e.g., form PTO/SB/08) of all patents, publications, or other information submitted for consideration by the office, either incorporated into this IDS or as an attachment hereto. A copy of each document listed is attached, except as explained below.

(check boxes A, B and/or C and fill in blanks, if appropriate.)

A. Pursuant to the Notice issued by the United States Patent and Trademark Office dated July 11, 2003 waiving the requirements of 37 C.F.R. § 1.98(a)(2)(i), a copy/copies of the United States Patent on PTO/SB08 is/are not being submitted.
B. Document(s) is (are) deemed substantially cumulative to document(s), and, in accordance with 1.98(c), only a copy of each of the latter documents is enclosed.
C. Certain documents were previously cited by or submitted to the Office in the following prior applications, which are relied upon under 35 U.S.C. 120:
< <insert &="" date="" filing="" no.="" serial="">&gt;</insert>
Applicant identifies these documents by attaching hereto copies of the forms PTO-892, PTO-14 and/or PTO/SB/08 from the files of the prior application(s) or a fresh PTO/SB/08 listing the documents, and request that they be considered and made of record in accordance with 1.98(d). 37 CFR 1.98(d), copies of these documents need not be filed in this application.
3. Cite No(s) are not in the English language. In accordance with 1.98(c), Applicant states:
An English translation of each document (or of the pertinent portions thereof), or a copy of each corresponding English-language patent or application, or English-language abstract (or claim) is enclosed.
The requirement for a concise explanation of the relevance of any foreign language document is satisfied by the attached search report; citation of the documents cited in the search report shall not be construed as an admission that they are or are considered to be, material to patentability of the subject matter claimed herein (See MPEP 860e).
A concise explanation of the relevance of document(s) is set forth as follows: [Insert concise explanation of
A concise explanation of the relevance of document(s) can be found on page(s) of the specification.
A concise explanation of document(s) can be found on the attached sheet.

application.]

_	English language (see reply to Comments 67 in the preamble to the final rules; 1135 OG 13 at 20).				
∐3.	Other information being provided for the consideration follows:	examiner's			
[A/An	Search Report, dated	, which issued during the			
prosecution of	Application No. which	corresponds to the present			

6. In accordance with 37 C.F.R. 1.97(g) and (h), the filing of this IDS should not be construed as a representation that a search has been made or that information cited is, or is considered to be, material to patentability as defined in §1.56 (b), or that any cited document listed or attached is (or constitutes) prior art. Unless other-wise indicated, the date of publication indicated for an item is taken from the face of the item and Applicant reserves the right to prove that the date of publication is in fact different.

Early and favorable consideration is earnestly solicited.

Our check in the amount of \$180.00 covering the fee set forth in 37 CFR 1.17(p) is enclosed. The Commissioner is authorized to charge any deficiency of up to \$300,00 or credit any excess in this fee to Denosit Account No. 04-0100.

Dated: September 12, 2006

Respectfully submitted.

Andrew K. Holmes Registration No.: 51,813 DARBY & DARBY P.C.

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Attorneys/Agents For Applicant

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Substitute for form 1449A/B/PTO	Complete If Known		$\neg$		
	Application Number	10/008,062-Conf. #5735			
INFORMATION DISCLOSURE	Filing Date	October 19, 2001	7		
STATEMENT BY APPLICANT	First Nemed Inventor	David A. Rizzieri			

(Use as many sheets as nec af

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U.S. PATENT DOCUMENTS					
Examiner Initials*	Cite No.1	Document Number  Number-Kind Code <sup>2</sup> (Finown)	Publication Date MM-DD-YYYY	Name of Palentee or Applicant of Cited Document	Pages, Columns, Lines, Whon Rolevent Passages or Relevan Figures Appear

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FOREIGN PATENT DOCUMENTS						
Examiner Initials*	Cite No.1	Foreign Patent Document Country Code*-Number*-Hind Code* (Filmown)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Otted Document	Pages, Columns, Lines. Where Relevent Pessages or Relevant Figures Appear	T <sup>0</sup>

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NON PATENT LITERATURE DOCUMENTS				
Examiner Initials	No. magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.		T <sup>1</sup>	
	CA	Zalutsky et al., Pharmacokinetics and tumor localization of "2" Hebaled anti-tanascin monocional antibody 81 C6 in patients with gillomes and other intracranial malignancies, Cancar Res., vol. 49, pp. 2807-2813 (May 15, 1989)		
	CB	Zalutsky et al., Monocionel antibody and F(eb) <sub>2</sub> fragment delivery to turnor in patients with glioma: comparison of intracarolid and intravanous administration, <u>Cancer Res.</u> , vol. 50, pp. 4105-4110, Livly 1, 1990.		
	СС	Rizziori et al., Phase I Trial study of *** Habeled chimoric 81 CS monoclonal antibody for tha treatment of patients with non-Hodykin's lymphoma, Clinical Observations, Interventions, And Thorspectic Trials, vol. 104, no. 3, no. 642-648 (Jupust 1, 2004).		

"EXAMINER, Initial if reference considered, whether or not diction is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

'Applicant's unique citation designation number (optional). "Applicant is to place a check mark have if English language Translaton is attached.

Examiner	Date
Signature	Considered